

REC'D 30 APR 2004

WIPO

PCT



Bescheinigung

Certificate

Attestation

Die angehefteten Unterlagen stimmen mit der ursprünglich eingereichten Fassung der auf dem nächsten Blatt bezeichneten internationalen Patentanmeldung überein.

The attached documents are exact copies of the international patent application described on the following page, as originally filed.

Les documents fixés à cette attestation sont conformes à la version initialement déposée de la demande de brevet international spécifiée à la page suivante.

BEST AVAILABLE COPY

Den Haag, den
The Hague,
La Haye, le

21 04 2004

Der Präsident des Europäischen Patentamts
Im Auftrag
For the President of the European Patent Office
Le Président de l'Office européen des brevets
p. o.


Mme. C. Rossi

**PRIORITY
DOCUMENT**

SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

Patentanmeldung Nr. PCT/EP 03/00482
Patent application no.
Demande de brevet n°

15-01-04

Blatt 2 der Bescheinigung
Sheet 2 of the certificate
Page 2 de l'attestation



Anmeldung Nr.:
 Application no.:
 Demande n°:

PCT/EP 03/00482

Anmelder:
 Applicant(s):
 Demandeur(s):

1. PIRELLI & C. S.p.A - Milano, Italy
2. PEREGO, Gabriele - Milano, Italy (US only)
3. SCENZA, Cristiana - Milano, Italy (US only)

Bezeichnung der Erfindung:
 Title of the invention:
 Titre de l'invention:

Cable with recyclable covering layer

Anmeldetag:
 Date of filing:
 Date de dépôt:

20 January 2003 (12.01.2003)

In Anspruch genommene Priorität(en)
 Priority(ies) claimed
 Priorité(s) revendiquée(s)

Staat:
 State:
 Pays:

Tag:
 Date:
 Date:

Aktenzeichen:
 File no.
 Numéro de dépôt:

Benennung von Vertragsstaaten : Siehe Formblatt PCT/RO/101 (beigefügt)
 Designation of contracting states : See Form PCT/RO/101 (enclosed)
 Désignation d'états contractants : Voir Formulaire PCT/RO/101 (ci-joint)

Bemerkungen:
 Remarks:
 Remarques:

Further applicants:

4. DELL'ANNA, Gaia - Milano, Italy (US only)
5. BELLI, Sergio - Milano, Italy (US only)

The applicant's name at the time of filing of the application was as follows:
 PIRELLI S.p.A

The registration of the change has taken effect on 27 November 2003 (27.11.2003)

Sheet No. ...3...

Box No. V DESIGNATION OF STATES

Mark the applicable check-boxes below; at least one must be marked.

The following designations are hereby made under Rule 4.9(a):

Regional Patent

- ☒ **AP** ARIPO Patent: GH Ghana, GM Gambia, KE Kenya, LS Lesotho, MW Malawi, MZ Mozambique, SD Sudan, SL Sierra Leone, SZ Swaziland, TZ United Republic of Tanzania, UG Uganda, ZM Zambia, ZW Zimbabwe, and any other State which is a Contracting State of the Harare Protocol and of the PCT (if other kind of protection or treatment desired, specify on dotted line)
- ☒ **EA** Eurasian Patent: AM Armenia, AZ Azerbaijan, BY Belarus, KG Kyrgyzstan, KZ Kazakhstan, MD Republic of Moldova, RU Russian Federation, TJ Tajikistan, TM Turkmenistan, and any other State which is a Contracting State of the Eurasian Patent Convention and of the PCT
- ☒ **EP** European Patent: AT Austria, BE Belgium, CH & LI Switzerland and Liechtenstein, CY Cyprus, DE Germany, DK Denmark, ES Spain, FI Finland, FR France, GB United Kingdom, GR Greece, IE Ireland, IT Italy, LU Luxembourg, MC Monaco, NL Netherlands, PT Portugal, SE Sweden, TR Turkey, and any other State which is a Contracting State of the European Patent Convention and of the PCT
- ☒ **OA** OAPI Patent: BF Burkina Faso, BJ Benin, CF Central African Republic, CG Congo, CI Côte d'Ivoire, CM Cameroon, GA Gabon, GN Guinea, GQ Equatorial Guinea, GW Guinea-Bissau, ML Mali, MR Mauritania, NE Niger, SN Senegal, TD Chad, TG Togo, and any other State which is a member State of OAPI and a Contracting State of the PCT (if other kind of protection or treatment desired, specify on dotted line)

National Patent (if other kind of protection or treatment desired, specify on dotted line):

- | | | |
|--|---|---|
| <input checked="" type="checkbox"/> AE United Arab Emirates | <input checked="" type="checkbox"/> GM Gambia | <input checked="" type="checkbox"/> NZ New Zealand |
| <input checked="" type="checkbox"/> AG Antigua and Barbuda | <input checked="" type="checkbox"/> HR Croatia | <input checked="" type="checkbox"/> OM Oman |
| <input checked="" type="checkbox"/> AL Albania | <input checked="" type="checkbox"/> HU Hungary | <input checked="" type="checkbox"/> PH Philippines |
| <input checked="" type="checkbox"/> AM Armenia | <input checked="" type="checkbox"/> ID Indonesia | <input checked="" type="checkbox"/> PL Poland |
| <input checked="" type="checkbox"/> AT Austria | <input checked="" type="checkbox"/> IL Israel | <input checked="" type="checkbox"/> PT Portugal |
| <input checked="" type="checkbox"/> AU Australia | <input checked="" type="checkbox"/> IN India | <input checked="" type="checkbox"/> RO Romania |
| <input checked="" type="checkbox"/> AZ Azerbaijan | <input checked="" type="checkbox"/> IS Iceland | <input checked="" type="checkbox"/> RU Russian Federation |
| <input checked="" type="checkbox"/> BA Bosnia and Herzegovina | <input checked="" type="checkbox"/> JP Japan | |
| <input checked="" type="checkbox"/> BB Barbados | <input checked="" type="checkbox"/> KE Kenya | <input checked="" type="checkbox"/> SD Sudan |
| <input checked="" type="checkbox"/> BG Bulgaria | <input checked="" type="checkbox"/> KG Kyrgyzstan | <input checked="" type="checkbox"/> SE Sweden |
| <input checked="" type="checkbox"/> BR Brazil | <input checked="" type="checkbox"/> KP Democratic People's Republic of Korea | <input checked="" type="checkbox"/> SG Singapore |
| <input checked="" type="checkbox"/> BY Belarus | <input checked="" type="checkbox"/> KR Republic of Korea | <input checked="" type="checkbox"/> SI Slovenia |
| <input checked="" type="checkbox"/> BZ Belize | <input checked="" type="checkbox"/> KZ Kazakhstan | <input checked="" type="checkbox"/> SK Slovakia |
| <input checked="" type="checkbox"/> CA Canada | <input checked="" type="checkbox"/> LC Saint Lucia | <input checked="" type="checkbox"/> SL Sierra Leone |
| <input checked="" type="checkbox"/> CH & LI Switzerland and Liechtenstein | <input checked="" type="checkbox"/> LK Sri Lanka | <input checked="" type="checkbox"/> TJ Tajikistan |
| <input checked="" type="checkbox"/> CN China | <input checked="" type="checkbox"/> LR Liberia | <input checked="" type="checkbox"/> TM Turkmenistan |
| <input checked="" type="checkbox"/> CO Colombia | <input checked="" type="checkbox"/> LS Lesotho | <input checked="" type="checkbox"/> TN Tunisia |
| <input checked="" type="checkbox"/> CR Costa Rica | <input checked="" type="checkbox"/> LT Lithuania | <input checked="" type="checkbox"/> TR Turkey |
| <input checked="" type="checkbox"/> CU Cuba | <input checked="" type="checkbox"/> LU Luxembourg | <input checked="" type="checkbox"/> TT Trinidad and Tobago |
| <input checked="" type="checkbox"/> CZ Czech Republic | <input checked="" type="checkbox"/> LV Latvia | |
| <input checked="" type="checkbox"/> DE Germany | <input checked="" type="checkbox"/> MA Morocco | <input checked="" type="checkbox"/> TZ United Republic of Tanzania |
| <input checked="" type="checkbox"/> DK Denmark | <input checked="" type="checkbox"/> MD Republic of Moldova | <input checked="" type="checkbox"/> UA Ukraine |
| <input checked="" type="checkbox"/> DM Dominica | <input checked="" type="checkbox"/> MG Madagascar | <input checked="" type="checkbox"/> UG Uganda |
| <input checked="" type="checkbox"/> DZ Algeria | <input checked="" type="checkbox"/> MK The former Yugoslav Republic of Macedonia | <input checked="" type="checkbox"/> US United States of America |
| <input checked="" type="checkbox"/> EC Ecuador | <input checked="" type="checkbox"/> MIN Mongolia | <input checked="" type="checkbox"/> UZ Uzbekistan |
| <input checked="" type="checkbox"/> EE Estonia | <input checked="" type="checkbox"/> MW Malawi | <input checked="" type="checkbox"/> VN Viet Nam |
| <input checked="" type="checkbox"/> ES Spain | <input checked="" type="checkbox"/> MX Mexico | <input checked="" type="checkbox"/> YU Yugoslavia |
| <input checked="" type="checkbox"/> FI Finland | <input checked="" type="checkbox"/> MZ Mozambique | <input checked="" type="checkbox"/> ZA South Africa |
| <input checked="" type="checkbox"/> GB United Kingdom | <input checked="" type="checkbox"/> NO Norway | <input checked="" type="checkbox"/> ZM Zambia |
| <input checked="" type="checkbox"/> GD Grenada | | <input checked="" type="checkbox"/> ZW Zimbabwe |
| <input checked="" type="checkbox"/> GE Georgia | | |
| <input checked="" type="checkbox"/> GH Ghana | | |

Check-boxes below reserved for designating States which have become party to the PCT after issuance of this sheet:

<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Precautionary Designation Statement: In addition to the designations made above, the applicant also makes under Rule 4.9(b) all other designations which would be permitted under the PCT except any designation(s) indicated in the Supplemental Box as being excluded from the scope of this statement. The applicant declares that those additional designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be regarded as withdrawn by the applicant at the expiration of that time limit. (Confirmation (including fees) must reach the receiving Office within the 15-month time limit.)

1.

CABLE WITH RECYCLABLE COVERING LAYER

The present invention relates to a cable with recyclable covering layer. In particular, the invention relates to a cable for transporting or distributing medium or high voltage electric energy, wherein an extruded covering layer based on a thermoplastic polymer material in admixture with a dielectric liquid with good mechanical and electrical properties is present, enabling, in particular, the use of high operating temperatures and the transportation of high power energy.

Said cable may be used for both direct current (DC) or alternating current (AC) transmission or distribution.

The requirement for products of high environmental compatibility, composed of materials which, in addition to not being harmful to the environment during production or utilization, may be easily recycled at the end of their life, is now fully accepted in the field of electrical and telecommunications cables.

However the use of materials compatible with the environment is conditioned by the need to limit costs while, for the more common uses, providing a performance equal to or better than that of conventional materials.

In the case of cables for transporting medium and high voltage energy, the various coverings surrounding the conductor commonly consist of polyolefin-based crosslinked polymer, in particular crosslinked polyethylene (XLPE), or elastomeric ethylene/propylene (EPR) or ethylene/propylene/diene (EPDM) copolymers, also crosslinked. The crosslinking, effected after the step of extrusion of the polymeric material onto the conductor, gives

31030

2

the material satisfactory mechanical and electrical properties even under high temperatures both during continuous use and with current overload.

It is well known however that crosslinked materials cannot be recycled, so that manufacturing wastes and the covering material of cables which have reached the end of their life may be disposed of only by incineration.

Electric cables are also known having their insulation consisting of a multi-layer wrapping of a paper or paper/polypropylene laminate impregnated with a large quantity of a dielectric liquid (commonly known as mass impregnated cables or also oil-filled cables). By completely filling the spaces present in the multi-layer wrapping, the dielectric liquid prevents partial discharges arising with consequent break down of the electrical insulation. As dielectric liquids products are commonly used such as mineral oils, polybutenes, alkylbenzenes and the like (see, for example, US 4,543,207, US 4,621,302, EP 987,718, WO 98/32137).

It is however well known that mass impregnated cables have numerous drawbacks compared with extruded insulation cables, so that their use is currently restricted to specific fields of application, in particular to the construction of high and very high voltage direct current transmission lines, both for terrestrial and in particular for underwater installations. In this respect, the production of mass impregnated cables is particularly complex and costly, both for the high cost of the laminates and for the difficulties encountered during the steps of wrapping the laminate and then of impregnating it with the dielectric liquid. In particular, the dielectric liquid used must have low viscosity under low

31030

3

temperatures to allow rapid and uniform impregnation, while at the same time it must have a low tendency to migrate during installation and operation of the cable to prevent liquid loss from the cable ends or from accidentally breaks on the cable. In addition, mass impregnated cables cannot be recycled and their use is limited to an operating temperature of less than 90°C.

Within non-crosslinked polymeric materials, it is known to use high density polyethylene (HDPE) for covering high voltage cables. HDPE has however the drawback of a lower temperature resistance than XLPE, both to current overload and during operation.

Thermoplastic low density polyethylene (LDPE) insulating coverings are also used in medium and high voltage cables: again in this case, these coverings are limited by a too low operating temperature (about 70°C).

International Patent Application WO 99/13477 discloses an insulating material consisting of a thermoplastic polymer forming a continuous phase which incorporates a liquid or easily meltable dielectric forming a mobile interpenetrating phase within the solid polymer structure. The weight ratio of thermoplastic polymer to dielectric is between 95:5 and 25:75. The insulating material may be produced by mixing the two components while hot either batchwise or continuously (for example, by means of an extruder). The resultant mixture is then granulated and used as insulating material for producing a high voltage electric cable by extrusion onto a conductor. The material may be used either in thermoplastic or crosslinked form. As thermoplastic polymers are indicated: polyolefins, polyacetates, cellulose polymers, polyesters, polyketones, polyacrylates, polyamides and polyamines. The use of

1030

4

polymers of low crystallinity is particularly suggested. The dielectric is preferably a synthetic or mineral oil of low or high viscosity, in particular a polyisobutene, naphthene, polyaromatic, 5 α -olefin or silicone oil.

International Patent Application WO 02/03398 in the name of the Applicant, discloses a cable comprising at least one electrical conductor and at least one extruded covering layer based on 10 thermoplastic polymer material in admixture with a dielectric liquid, wherein said thermoplastic material comprises a propylene homopolymer or a copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin 15 other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 140°C and a melting enthalpy of from 30 J/g to 100 J/g. Said dielectric liquid comprises at least one alkylaryl hydrocarbon having at least two non- 20 condensed aromatic rings and a ratio of number of aryl carbon atoms to total number of carbon atoms greater than or equal to 0.6, preferably greater than or equal to 0.7.

International Patent Application WO 02/27731 in 25 the name of the Applicant, discloses a cable comprising at least one electrical conductor and at least one extruded covering layer based on thermoplastic polymer material in admixture with a dielectric liquid, wherein said thermoplastic 30 material comprises a propylene homopolymer or a copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 35 140°C and a melting enthalpy of from 30 J/g to 100 J/g. Said dielectric liquid comprises at least one

C1030

5

diphenyl ether, non-substituted or substituted with at least one linear or branched, aliphatic, aromatic or mixed aliphatic and aromatic C₁-C₃₀ hydrocarbon radical.

5 However, the prior art above cited presents some drawbacks.

10 As a matter of fact, Applicant noticed that the addition of a dielectric liquid to a polymer material should both determine a significant
15 increase in its electrical properties (in particular, its dielectric strength), without impairing its thermomechanical characteristics and without resulting in exudation of the dielectric liquid from the polymer material. In particular, the
20 resultant cable should give substantially constant mechanical and electrical performances with time and hence high reliability, even at high operating temperatures (at least 90°C and beyond, in particular at operating temperature up to 110°C for
25 continuous use and up to 140°C in the case of current overload). In particular, Applicant noticed that the presence of two phases, e.g. a continuous phase of a thermoplastic material and an additional phase incorporated therein of a dielectric liquid,
30 with the consequent microscopically non homogeneous dispersion of said dielectric liquid onto said thermoplastic material, does not allow to obtain all the above reported characteristics.

35 The Applicant has now found that it possible to overcome said drawbacks by using, as recyclable polymer base material, at least one thermoplastic propylene homopolymer or copolymer or a mechanical mixture of said at least one thermoplastic propylene homopolymer or copolymer with at least one elastomeric copolymer of ethylene with at least one aliphatic α -olefin, and optionally a polyene, mixed

PC1030

6

with at least one dielectric liquid as hereinafter defined. The resultant composition possesses suitable flexibility, excellent thermomechanical characteristics and high electrical performance, such as to make it particularly suitable for forming at least one covering layer, and in particular an electrical insulating layer, of a medium or high voltage cable of high operating temperature, of at least 90°C and beyond, in particular at operating temperature up to 110°C for continuous use and up to 140°C in the case of current overload. The dielectric liquid suitable for implementing the invention has high compatibility with the polymer base material and high efficiency in the sense of improving electrical performance, consequently allowing the use of small quantities (e.g. quantities not higher than the saturation concentration of the dielectric liquid in the polymer base material) of said dielectric liquid such as not to impair the thermomechanical characteristics of the insulating layer and to avoid the exudation of said dielectric liquid from the polymer base material.

High compatibility between the dielectric liquid and the polymer base material allows to obtain a microscopically homogeneous dispersion of the dielectric liquid in the polymer base material. Moreover, the dielectric liquid suitable for forming the cable covering layer of the present invention comprises a small quantity of polar compounds, in order to avoid a significant increasing of the dielectric losses. It has to be noticed also that the use of a dielectric liquid with a relatively low melting point (e.g. a melting point not higher than 80°C) does not give rise to manufacturing problems both during the mixing with the polymer material and during the production of the cable. As a matter of

01030

7

fact, the low melting point allows to an easier handling of the dielectric liquid which may be easily melted without the need of additional and complex manufacturing steps (e.g., a melting step of the dielectric liquid) and/or apparatuses. Moreover, Applicant noticed also that, when dielectric liquid is aromatic, high compatibility with the polymer base material may be achieved even in the presence of dielectric liquid with a low ratio of number of aromatic carbon atoms to total number of carbon atoms (e.g., ratio lower than 0.6).

The Applicant has also noticed that the addition of said dielectric liquid reduces or even eliminates the optical phenomena commonly known as "stress whitening" thanks to the fact that said dielectric liquid is microscopically homogeneously dispersed in the polymer material.

According to a first aspect, the present invention relates to a cable comprising at least one electrical conductor and at least one extruded covering layer based on a thermoplastic polymer material in admixture with a dielectric liquid, wherein:

- said thermoplastic polymer material is selected from:

(a) at least one propylene homopolymer or at least one copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 130°C and a melting enthalpy of from 20 J/g to 100 J/g;

(b) a mechanical mixture comprising at least one propylene homopolymer or copolymer (a) and

(c) at least one elastomeric copolymer of

31030

8

ethylene with at least one aliphatic α -olefin, and optionally a polyene;

- the concentration by weight of said dielectric liquid in said thermoplastic polymer material is less or equal to the saturation concentration of said dielectric liquid in said thermoplastic polymer material;
- said dielectric liquid has the following characteristics:
 - an amount of polar compounds lower than or equal to 2.5% by weight with respect to the total weight of the dielectric liquid;
 - a melting point or a pour point lower than 80°C;
 - a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms lower than 0.6, when the dielectric liquid is aromatic.

In the present description and in the subsequent claims, the term "conductor" means a conducting element as such, of elongated shape and preferably of a metallic material, or a conducting element coated with a semiconducting layer.

The saturation concentration of the dielectric liquid in the thermoplastic polymer material may be determined by a liquid absorption method on Dumbell samples: further details regarding said method will be described in the examples given hereinbelow.

The amount of polar compounds of the dielectric liquid may be determined according to ASTM standard D2007-02.

The melting point may be determined by known techniques such as, for example, by Differential Scanning Calorimetry (DSC) analysis.

The pour point may be determined according to ASTM standard D97.

11030

9

The ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms may be determined according to ASTM standard D3238.

5 According to a first embodiment, the extruded covering layer based on said thermoplastic polymer material in admixture with said dielectric liquid is an electrically insulating layer.

10 According to a further embodiment, the extruded covering layer based on said thermoplastic polymer material in admixture with said dielectric liquid is a semiconductive layer.

15 According to one preferred embodiment, the propylene homopolymer or copolymer (a) which may be used in the present invention has a melting point of from 140°C to 170°C.

Preferably, the propylene homopolymer or copolymer (a) has a melting enthalpy of from 30 J/g to 85 J/g.

20 Said melting enthalpy (ΔH_m) may be determined by Differential Scanning Calorimetry (DSC) analysis.

Preferably, the propylene homopolymer or copolymer (a) has a flexural modulus, measured according to ASTM standard D790, at room temperature, of from 30 MPa to 1400 MPa, and more preferably from 60 MPa to 1000 MPa.

25 Preferably, the propylene homopolymer or copolymer (a) has a melt flow index (MFI), measured at 230°C with a load of 21.5 N according to ASTM standard D1238/L, of from 0.05 dg/min to 10.0 dg/min, more preferably from 0.4 dg/min to 5.0 dg/min.

35 If a copolymer of propylene with at least one olefin comonomer (a) is used, this latter is preferably present in a quantity of less than or equal to 15 mol%, and more preferably of less than or equal to 10 mol%. The olefin comonomer is, in

PC1030

10

particular, ethylene or an α -olefin of formula $\text{CH}_2=\text{CH-R}$, where R is a linear or branched $\text{C}_2\text{-C}_{10}$ alkyl, selected, for example, from: 1-butene, 1-pentene, 4-methyl-1-pentene, 1-hexene, 1-octene, 1-decene, 1-dodecene, or mixtures thereof. Propylene/ethylene copolymers are particularly preferred.

Preferably, said propylene homopolymer or copolymer (a) is selected from:

- 10 (a₁) a propylene homopolymer or a copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, having a flexural modulus generally of from 30 MPa to 900 MPa, and
- 15 preferably of from 50 MPa to 400 MPa;
- (a₂) a heterophase copolymer comprising a thermoplastic phase based on propylene and an elastomeric phase based on ethylene
- 20 copolymerized with an α -olefin, preferably with propylene, in which the elastomeric phase is preferably present in a quantity of at least 45 wt% with respect to the total weight of the heterophase copolymer.

Particularly preferred of said class (a₁) is a

25 propylene homopolymer or a copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having:

- a melting point of from 140°C to 170°C;
- 30 - a melting enthalpy of from 30 J/g to 80 J/g;
- a fraction soluble in boiling diethyl ether in an amount of less than or equal to 12 wt%, preferably from 1 wt% to 10 wt%, having a melting enthalpy of less than or equal to 4 J/g,
- 35 - preferably less than or equal to 2 J/g;
- a fraction soluble in boiling n-heptane in an

PC1030

11

amount of from 15 wt% to 50 wt%, preferably from 20 wt% to 50 wt%, having a melting enthalpy of from 10 J/g to 40 J/g, preferably from 15 J/g to 30 J/g; and

- 5 - a fraction insoluble in boiling n-heptane in an amount of from 40 wt% to 85 wt%, preferably from 50 wt% to 80 wt%, having a melting enthalpy of greater than or equal to 45 J/g, preferably from 50 J/g to 95 J/g.

- 10 Further details concerning these materials and their use in cables covering are given in International Patent Application WO 01/37289 in the name of the Applicant.

- The heterophase copolymers of class (a₂) are
15 obtained by sequential copolymerization of: i) propylene, possibly containing minor quantities of at least one olefin comonomer selected from ethylene and an α -olefin other than propylene; and then of:
ii) a mixture of ethylene with an α -olefin, in
20 particular propylene, and possibly with minor portions of a diene.

- Particularly preferred of said class (a₂) is a heterophase copolymer in which the elastomeric phase consists of an elastomeric copolymer of ethylene and
25 propylene comprising from 15 wt% to 50 wt% of ethylene and from 50 wt% to 85 wt% of propylene with respect to the weight of the elastomeric phase. Further details concerning these materials and their use in cables covering are given in International
30 Patent Application WO 00/41187 in the name of the Applicant.

- Products of class (a₁) are available commercially for example under the trademark Rexflex[®] WL 105 of Huntsman Polymer Corporation or Borsoft[®] SA 233 CF
35 of Borealis.

Products of class (a₂) are available commercially

C1030

12

for example under the trademark Hifax[®] CA 10 A, Moplen[®] EP 310 G, or Adflex[®] Q 200 F of Basell.

According to one preferred embodiment, the elastomeric copolymer of ethylene (c) has a melting
5 enthalpy of less than 30 J/g. The quantity of said elastomeric copolymer (c) is generally less than 70% by weight, preferably of from 20% by weight to 60% by weight, with respect to the total weight of the thermoplastic base material.

10 With reference to the elastomeric copolymer of ethylene (c), the term "aliphatic α -olefin" generally means an olefin of formula $\text{CH}_2=\text{CH-R}$, in which R represents a linear or branched alkyl group containing from 1 to 12 carbon atoms. Preferably,
15 the aliphatic α -olefin is selected from propylene, 1-butene, isobutylene, 1-pentene, 4-methyl-1-pentene, 1-hexene, 1-octene, 1-dodecene, or mixtures thereof. Propylene, 1-hexene and 1-octene are particularly preferred.

20 With reference to the elastomeric copolymer of ethylene (c), the term "polyene" generally means a conjugated or non-conjugated diene, triene or tetraene. When a diene comonomer is present, this comonomer generally contains from 4 to 20 carbon
25 atoms and is preferably selected from: linear conjugated or non-conjugated diolefins such as, for example, 1,3-butadiene, 1,4-hexadiene, 1,6-octadiene, and the like; monocyclic or polycyclic dienes such as, for example, 1,4-cyclohexadiene,
30 5-ethylidene-2-norbornene, 5-methylene-2-norbornene, vinylnorbornene, or mixtures thereof. When a triene or tetraene comonomer is present, this comonomer generally contains from 9 to 30 carbon atoms and is preferably selected from trienes or tetraenes
35 containing a vinyl group in the molecule or a 5-norbornen-2-yl group in the molecule. Specific

PC1030

13

examples of triene or tetraene comonomers which may be used in the present invention are: 6,10-dimethyl-1,5,9-undecatriene, 5,9-dimethyl-1,4,8-decatriene, 6,9-dimethyl-1,5,8-decatriene, 6,8,9-trimethyl-1,6,8-decatriene, 6,10,14-trimethyl-1,5,9,13-pentadecatetraene, or mixtures thereof. Preferably, the polyene is a diene.

Particularly preferred elastomeric copolymers of ethylene (c) are:

10 (c₁) copolymers having the following monomer composition: 35 mol%-90 mol% of ethylene; 10 mol%-65 mol% of an aliphatic α -olefin, preferably propylene; 0 mol%-10 mol% of a polyene, preferably a diene, more preferably, 15 1,4-hexadiene or 5-ethylene-2-norbornene (EPR and EPDM rubbers belong to this class);

(c₂) copolymers having the following monomer composition: 75 mol%-97 mol%, preferably 90 mol%-95 mol%, of ethylene; 3 mol%-25 mol%, preferably 5 mol%-10 mol%, of an aliphatic α -olefin; 0 mol%-5 mol%, preferably 0 mol%-2 mol%, of a polyene, preferably a diene (for example ethylene/1-octene copolymers, such as the products Engage[®] of DuPont-Dow Elastomers).

25 According to a preferred embodiment, the dielectric liquid has an amount of polar compounds of between 0.1 and 2.3.

According to a further preferred embodiment, the dielectric liquid has a melting point or a pour point of between -130°C and +80°C.

30 According to a further preferred embodiment, the dielectric liquid has a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of between 0.01 and 0.4.

35 According to a further preferred embodiment, the dielectric liquid preferably has a dielectric

.030

14

constant, at 25°C, of less than or equal to 3.5 and preferably less than 3 (measured in accordance with IEC 247).

According to a further preferred embodiment, the dielectric liquid has a predetermined viscosity in order to prevent fast diffusion of the liquid within the insulating layer and hence its outward migration, as well as to enable the dielectric liquid to be easily fed and mixed into the thermoplastic polymer material. Generally, the dielectric liquid of the invention has a viscosity, at 40°C, of between 10 cst and 800 cst, preferably between 20 cst and 500 cst (measured according to ASTM standard D445).

According to one preferred embodiment, the dielectric liquid may be selected from: mineral oils such as, for example, naphthenic oils, aromatic oils, paraffinic oils, polyaromatic oils, said mineral oils optionally containing at least one heteroatom selected from oxygen, nitrogen or sulphur; liquid paraffins; vegetable oils such as, for example, soybean oil, linseed oil, castor oil; oligomeric aromatic polyolefins; paraffinic waxes such as, for example, polyethylene waxes, polypropylene waxes; synthetic oils such as, for example, silicone oils, alkyl benzenes (such as, for example, dodecylbenzene, di(octylbenzyl)toluene), aliphatic esters (such as, for example, tetraesters of pentaerythritol, esters of sebacic acid, phthalic esters), olefin oligomers (such as, for example, optionally hydrogenated polybutenes or polyisobutenes), or mixtures thereof. Paraffinic oils and naphthenic oils are particularly preferred.

The dielectric liquid suitable for implementing the invention has good heat resistance, considerable gas absorption capacity, in particular hydrogen

C1030

15

absorption, and high resistance to partial discharges, so that dielectric losses are limited even at high temperature and high electrical gradient. The weight ratio of dielectric liquid to thermoplastic polymer material of the present invention is generally between 1:99 and 25:75, preferably between 2:98 and 20:80, and more preferably between 3:97 and 10:90.

According to one preferred embodiment, the cable of the invention has at least one extruded covering layer with electrical insulation properties formed from the thermoplastic polymer material in admixture with the aforescribed dielectric liquid.

According to a further preferred embodiment, the cable of the invention has at least one extruded covering layer with semiconductive properties formed from the thermoplastic polymer material in admixture with the aforescribed dielectric liquid. To form a semiconductive layer, a conductive filler is generally added to the polymer material. To ensure good dispersion of the conductive filler within the thermoplastic polymer material, the latter is preferably selected from propylene homopolymers or copolymers comprising at least 40 wt% of amorphous phase, with respect to the total polymer weight.

According to a further aspect, the present invention relates to a polymer composition comprising a thermoplastic polymer material in admixture with a dielectric liquid, wherein:

- said thermoplastic polymer material is selected from:

(a) at least one propylene homopolymer or at least one copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer

C1030

16

having a melting point greater than or equal to 130°C and a melting enthalpy of from 20 J/g to 100 J/g;

- 5 (b) a mechanical mixture comprising at least one propylene homopolymer or copolymer (a) and (c) at least one elastomeric copolymer of ethylene with at least one aliphatic α -olefin, and optionally a polyene;
- 10 - the concentration by weight of said dielectric liquid in said thermoplastic polymer material is less or equal to the saturation concentration of said dielectric liquid in said thermoplastic polymer material;
- 15 - said dielectric liquid has the following characteristics:
- an amount of polar compounds lower than or equal to 2.5% by weight with respect to the total weight of the dielectric liquid;
 - 20 - a melting point or a pour point lower than 80°C;
 - a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms lower than 0.6, when the dielectric liquid is aromatic.

25 According to a further aspect, the present invention relates to the use of a polymer composition, as described hereinabove, as the polymer base material for preparing a cable covering layer with electrical insulation properties, or for

30 preparing a cable covering layer with semiconductive properties.

In forming a covering layer for the cable of the invention, other conventional components may be added to the aforesaid polymer composition, such

35 as antioxidants, processing aids, water tree retardants, or mixtures thereof.

1030

17

Conventional antioxidants suitable for the purpose are for example distearyl- or dilauryl-thiopropionate and pentaerythrityl-tetrakis [3-(3,5-di-t-butyl-4-hydroxyphenyl)propionate], or mixtures thereof.

Processing aids which may be added to the polymer composition include, for example, calcium stearate, zinc stearate, stearic acid, or mixtures thereof.

With particular reference to medium and high voltage cables, the polymer materials as defined hereinabove may be advantageously used to obtain an insulating layer. As stated above, these polymer base materials show indeed good mechanical characteristics both at ambient temperature and under hot conditions, and also show improved electrical properties. In particular they enable high operating temperature to be reached, comparable with or even exceeding that of cables with coverings consisting of crosslinked polymer base materials.

If a semiconductive layer has to be formed, a conductive filler, in particular carbon black, is generally dispersed within the polymer base material in a quantity such as to provide the material with semiconductive characteristics (i.e. such as to obtain a resistivity of less than 5 Ohm*m at ambient temperature). This quantity is generally between 5 and 80 wt%, and preferably between 10 and 50 wt%, of the total weight of the mixture.

The use of the same polymer composition for both the insulating layer and the semiconductive layers is particularly advantageous in producing cables for medium or high voltage, in that it ensures excellent adhesion between adjacent layers and hence a good electrical behaviour, particularly at the interface between the insulating layer and the inner

C1030

18

semiconductive layer, where the electrical field and hence the risk of partial discharges are higher.

The polymer composition of the present invention may be prepared by mixing together the thermoplastic polymer material, the dielectric liquid and any other additives possibly present by using methods known in the art. Mixing may be carried out for example by an internal mixer of the type with tangential rotors (Banbury) or with interpenetrating rotors, or, preferably, in a continuous mixer of Kneader (Buss) type, or of co- or counter-rotating double-screw type.

Alternatively, the dielectric liquid of the present invention may be added to the thermoplastic polymer material during the extrusion step by direct injection into the extruder cylinder as disclosed, for example, in International Patent Application WO02/47092 in the name of the Applicant.

According to the present invention, the use of the aforedefined polymer composition in cable covering layers for medium or high voltage enables recyclable, flexible coverings to be obtained with excellent mechanical and electrical properties.

Greater compatibility has also been found between the dielectric liquid and the thermoplastic polymer material of the present invention than in the case of similar mixtures of the same polymer material with other dielectric liquids known in the art. This greater compatibility leads, *inter alia*, to less exudation of the dielectric liquid. Due to their high operating temperature and their low dielectric losses, the cables of the invention can carry, for the same voltage, a power at least equal to or even greater than that transportable by a traditional cable with XLPE covering.

For the purposes of the invention the term

1030

19

"medium voltage" generally means a voltage of between 1 kV and 35 kV, whereas "high voltage" means voltages higher than 35 kV.

Although this description is mainly focused on the production of cables for transporting or distributing medium or high voltage energy, the polymer composition of the invention may be used for covering electrical devices in general and in particular cables of different type, for example low voltage cables, telecommunications cables or combined energy/telecommunications cables, or accessories used in electrical lines, such as terminals, joints or connectors.

Further characteristics will be apparent from the detailed description given hereinafter with reference to the accompanying drawing, in which:

- Figure 1 is a perspective view of an electric cable, particularly suitable for medium or high voltage, according to the invention.

In Figure 1, the cable (1) comprises a conductor (2), an inner layer with semiconductive properties (3), an intermediate layer with insulating properties (4), an outer layer with semiconductive properties (5), a metal screen (6), and an outer sheath (7).

The conductor (2) generally consists of metal wires, preferably of copper or aluminium, stranded together by conventional methods, or of a solid aluminium or copper rod. At least one covering layer selected from the insulating layer (4) and the semiconductive layers (3) and (5) comprises the composition of the invention as heretofore defined. Around the outer semiconductive layer (5) there is usually positioned a screen (6), generally of electrically conducting wires or strips wound helically. This screen is then covered by a sheath

1030

20

(7) of a thermoplastic material such as, for example, non-crosslinked polyethylene (PE).

The cable can be also provided with a protective structure (not shown in Figure 1) the main purpose of which is to mechanically protect the cable against impacts or compressions. This protective structure may be, for example, a metal reinforcement or a layer of expanded polymer as described in WO 98/52197 in the name of the Applicant.

Figure 1 shows only one possible embodiment of a cable according to the invention. Suitable modifications known in the art can be made to this embodiment, but without departing from the scope of the invention.

The cable covering layer or layers of thermoplastic material according to the present invention may be manufactured in accordance with known methods, for example by extrusion. The extrusion is advantageously carried out in a single pass, for example by the tandem method in which individual extruders are arranged in series, or by co-extrusion with a multiple extrusion head.

The following examples illustrate the invention, but without limiting it.

EXAMPLES 1-5

Compositions preparation

The following components were used:

- a propylene heterophase copolymer with melting point 165°C, melting enthalpy 30 J/g, MFI 0.8 dg/min and flexural modulus 150 MPa (Adflex[®] Q 200 F - commercial product of Basell);
- a propylene heterophase copolymer with melting point 142°C, melting enthalpy 25 J/g, MFI 0.6 dg/min and flexural modulus of 85 MPa (Hifax[®] CA 10A - commercial product of Basell);
- Sunpar[®] 2280 (commercial product of Sunoco):

31030

21

- paraffinic oil with viscosity of 475 cSt at 40°C, pour point of -15°C and ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of 0.02, consisting of 69 %wt paraffinic carbon atoms, 29 wt% naphthenic carbon atoms, 2 wt% aromatic carbon atoms and 1.5 wt% polar compounds;
- 5 - Nyflex[®] 820 (commercial product of Nynas): naphthenic oil with viscosity of 110 cSt at 40°C, pour point of -27°C and ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of 0.1, consisting of 10 %wt aromatic carbon atoms, 46 wt% naphthenic carbon atoms, 44 wt% paraffinic carbon atoms and 0.2 wt% polar compounds;
- 10 15 - Nytex[®] 840 (commercial product of Nynas): naphthenic oil with with viscosity of 370 cSt at 40°C, pour point of -12°C and ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of 0.15, consisting of 15 %wt aromatic carbon atoms, 34 wt% naphthenic carbon atoms, 51 wt% paraffinic carbon atoms and 2.3 wt% polar compounds;.
- 20

The polymer in granular form was preheated, under agitation, at 80°C, over 15 min, in a turbomixer. Subsequently, the dielectric liquid, 6% by weight, was added to the preheated polymer. After the addition, agitation was continued for 2 hours at 80°C until the liquid was completely absorbed in the polymer granules.

25 30

After this first stage, the resultant material was kneaded in a laboratory double-screw Brabender Plasticorder PL2000 at a temperature of 180°C to complete homogenization. The resultant material left the double-screw mixer in the form of granules.

35

C1030

22

Measurement of dielectric losses

Plates of 0.5 mm thickness were formed from the material obtained as disclosed above. The plates were moulded at 195°C with 15 min preheating.

- 5 The plates obtained in this manner were subjected to dielectric loss measurement by measuring the tangent of the loss angle ($\tan\delta$) (according to ASTM standard D150) at various temperatures and at various gradients (G). The
- 10 measurements at $G = 10$ kV/mm were effected under a pressure of 25 bar of nitrogen. The $\tan\delta \times 10^{-4}$ ($G = 2$ kV/mm at 50 Hz) at 20°C was of between 3 and 50 and the $\tan\delta \times 10^{-4}$ ($G = 2$ kV/mm at 50 Hz) at 90°C was of between 5 and 100.

15 Measurement of flexural modulus

The flexural modulus was determined on plates 60 mm x 10 mm x 1.5 mm obtained as disclosed above in accordance with ASTM standard D790: the obtained results are given in Table 1.

20 Measurement of melting point (T_m) and melting enthalpy (ΔH)

- The melting point (T_m) and the melting enthalpy (ΔH) were determined by Differential Scanning Calorimetry (DSC) analysis by using a Mettler Toledo
- 25 DSC 820 differential scanning calorimeter. The temperature program below was applied to the sample to be analysed:

- cooling from room temperature to -100°C;
- heating from -100°C to 200°C at a rate of

30 10°C/min.;

- isotherm for 5 minutes at 200°C;
- cooling to -100°C at a rate of 2°C/min.;
- isotherm for 10 minutes at -100°C;
- heating to 200°C at a rate of 10°C/min.

35 The obtained results are given in Table 1.

C1030

23

TABLE 1

EXAMPLE	Flexural modulus (MPa)	Melting point (T _m) (°C)	Melting enthalpy (ΔH) (J/g)
1	37	162	40.2
2	35	163	40.9
3	30	160	41.1
4	60	139	30.7
5	60	140	32.0

Example 1: 94% by weight Adflex[®] Q 200 F + 6% by weight Sunpar[®] 2280;

5 Example 2: 94% by weight Adflex[®] Q 200 F + 6% by weight Nyflex[®] 820;

Example 3: 94% by weight Adflex[®] Q 200 F + 6% by weight Nyltex[®] 840;

10 Example 4: 94% by weight Hifax[®] CA 10 A + 6% by weight Sunpar[®] 2280;

Example 5: 94% by weight Hifax[®] CA 10 A + 6% by weight Nyltex[®] 2280.

EXAMPLE 6Measurement of the saturation concentration

15 In order to determine the saturation concentration of the dielectric liquid in the thermoplastic materials, a plurality of plates were manufactured starting from the raw materials in pellets.

20 Two plates (200 mm x 200 mm x 0.5 mm) were obtained by molding the raw material (Adflex[®] Q 200 F) at 190°C. Five smaller Dumbell samples were

obtained from each of the above plates and weighted (W_0).

The Dumbell samples were then totally immersed at 20°C, into a dielectric liquid: Sunpar[®] 2280 and Nyflex[®] 820, respectively. The saturation concentration was measured by determining the weight change (in percentage) of the plates after different times. The Dumbell samples were removed from the dielectric liquid after 3, 6, 9, 12 and 15 days, and after having cleaned their surface with a dry and clean cloth, they were weighted (W_1).

The dielectric liquid absorption was determined by the following formula:

$$\% \text{ of absorbed dielectric liquid} = [(W_1 - W_0)/W_0] \times 100.$$

The saturation concentration is reached when W_1 shows a variation lower than 1% with respect to the total weight increase which correspond to ($W_1 - W_0$).

The obtained results were the following:

- the saturation concentration of Sunpar[®] Q 200 F in the Adflex[®] Q 200 F is higher than 6% by weight;
- the saturation concentration of Nyflex[®] 820 in the Adflex[®] Q 200 F is higher than 12% by weight.

EXAMPLE 7

In order to verify the absence of two phases, e.g., the absence of a continuous phase of a thermoplastic material and of an additional phase incorporated therein of a dielectric liquid, samples of the dielectric liquid as such and of thermoplastic material additioned with the dielectric liquid were subjected to the Modulated Differential Scanning Calorimetry (MDSC) analysis using a TA Instrument DSC 2920 Modulated differential scanning calorimeter.

10 mg of each sample were subjected to the following temperature program:

C1030

25

- equilibrating at -145°C ;
 - modulating $\pm 0.48^{\circ}\text{C}$ every 60 seconds;
 - keeping at -145°C for 5 minutes;
 - heating to 200°C at a rate of $5^{\circ}\text{C}/\text{min}$;
 - 5 - keeping at 200°C for 2 minutes.
- The obtained results are given in Table 2.

TABLE 2

10

EXAMPLE	MDSC ANALYSIS
Sunpar [®] 2280	-0.59°C
Adflex [®] Q 200 F + 6% Sunpar [®] 2280	absent
Adflex [®] Q 200 F + 20% Sunpar [®] 2280	-0.59°C

The results in Table 2 show that:

- in the case the dielectric liquid as such, a peak at -0.59°C was present;
- 15 - in the case the dielectric liquid is added to the thermoplastic material in a quantity (6% by weight) not higher than its saturation concentration in said thermoplastic material, the peak at -0.59°C , characteristic of the dielectric liquid as such, was not present,
- 20 showing that the dielectric liquid was microscopically homogeneously dispersed in the thermoplastic material;
- in the case the dielectric liquid is added to the thermoplastic material in a quantity (20% by weight) higher than its saturation concentration in said thermoplastic material, the peak at
- 25 -0.59°C , characteristic of the dielectric liquid

1030

26

as such, was present, showing that the dielectric liquid was not microscopically homogeneously dispersed in the thermoplastic material.

5 EXAMPLES 8-9 (comparative).

Compositions preparation

The following components were used:

- a propylene heterophase copolymer with melting point of 142°C, melting enthalpy 25 J/g, melting point 142°C, MFI 0.6 dg/min and flexural modulus of 85 MPa (Hifax[®] CA 10A - commercial product of Basell);
- Nytex[®] 800 (commercial product of Nynas): naphthenic oil with viscosity of 7.3 cSt at 40°C, pour point of -60°C and ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of 0.07, consisting of 7 wt% aromatic carbon atoms, 53 wt% naphthenic carbon atoms, 40 wt% of paraffinic carbon atoms and 0.5 wt% polar compounds;
- Indopol[®] L-100 (commercial product of BP Amoco): polybutene oil with viscosity of 210 cSt at 40°C, pour point of -30°C and 0.5 wt% polar compounds.

25 The polymer in granular form was preheated, under agitation, at 80°C, over 15 min, in a turbomixer. Subsequently, the dielectric liquid, 6% by weight, was added to the preheated polymer. After the addition agitation was continued for 2 hours at 30 80°C until the liquid was completely absorbed in the polymer granules.

After this first stage, the resultant material was kneaded in a laboratory double-screw Brabender Plasticorder PL2000 at a temperature of 150°C to 35 complete homogenization. The resultant material left the double-screw mixer in the form of granules.

EXAMPLE 10Scanning Electron Microscopy (SEM)

Scanning Electron Microscopy (SEM) analysis was conducted as follows by utilizing the compositions of Examples 1-5 (according to the present invention) and the compositions of Examples 8-9 (comparative). Compression molded tensile samples were notched with a razor blade and subsequently immersed in liquid nitrogen. Samples were then fractured in a compact tension mode. Freeze-fracture morphology of gold coated samples was examined with a Hitachi S-400 SEM operating at 10 KV. Digital image analysis was performed on a series of micrographs to determine the presence of a single-phase material or of a two-phases material. At 5000X the surfaces of the samples obtained from the compositions of Examples 1-5 (according to the present invention) were homogeneous and devoid of cavity showing that the material is a single phase material. On the contrary, at 5000X, the surfaces of the samples obtained from the compositions of Example 6 and 7 (comparative), were not homogeneous and presented a lot of cavity showing that the material is a two phase material. Moreover, the samples obtained from Examples 8-9, showed exudation of the dielectric liquid at room temperature.

EXAMPLE 11Cable production

The compositions of the insulating layer and of the semiconductive layers are described in Table 3 below.

1030

28

5

TABLE 3

	Cable according to the present invention		Comparison cable	
	Inner and outer semicond. layers	Insulation layer	Inner and outer semicond. layers	Insulation layer
	(%) by weight	(%) by weight	(%) by weight	(%) by weight
Adflex [®] Q 200 F	60.4	93.4	66.4	99.4
Ensaco [®] 250 G	33	-	33	-
Sunpar [®] 2280	6	6	-	-
Irganox [®] PS 802	0.4	0.4	0.4	0.4
Irganox [®] 1010	0.2	0.2	0.2	0.2

Ensaco[®] 250 G: carbon black with specific surface of 65 m²/g (commercial product of MMM Carbon);

C1030

29

Irganox[®] PS 802 (antioxidant): distearyl
thiodipropionate (commercial product of Ciba
Specialty Chemicals);

5 Irganox[®] 1010 (antioxidant): pentaerithrityl-
tetrakis-(3-(3,5-di-t-butyl-4-hydroxy-phenyl)-
propionate (commercial product of Ciba Specialty
Chemicals).

The process used for manufacturing the cable was
the following.

10 The Adflex[®] Q 200 F was fed directly into the
extruder hopper. Subsequently, the Sunpar[®] 2280
previously mixed with the antioxidants, was injected
at high pressure into the extruder. An extruder
having a diameter of 80 mm and an L/D ratio of 25
15 was used. The injection was made during the
extrusion at about 20 D from the beginning of the
extruder screw by means of three injections point
on the same cross-section at 120° from each other.
The dielectric liquid was injected at a temperature
20 of 70°C and a pressure of 250 bar.

The cable leaving the extrusion head was cooled
to ambient temperature by passing it through cold
water.

25 The finished cable consisted of an aluminum
conductor (cross-section 150 mm²), an inner
semiconductive layer of about 0.5 mm in thickness,
an insulating layer of about 4.5 mm in thickness and
finally an outer semiconductive layer of about 0.5
mm in thickness.

30 Under similar conditions, by using the materials
indicated in Table 2, a comparison cable was
produced without adding the dielectric liquid.

Dielectric strength

35 Three pieces (each being 20 metres in length) of
the two cables produced as described above were
subjected to dielectric strength measurement using

1030

30

alternating current at ambient temperature. Starting from 100 kV the gradient applied to the cables was increased by 10 kV every 10 minutes until the cables broke down. The break down gradient considered is that on the conductor.

Table 4 summarizes the results of the electrical tests: the data represent the average value obtained from three different measurements.

TABLE 4

	Cable according to the present invention	Comparison cable
	(kV/mm)	(kV/mm)
AC break down	59	29

EXAMPLE 12 (comparison)Cable production

The compositions of the insulation layer is described in Table 5 below.

TABLE 5

COMPOSITION OF THE INSULATION LAYER	
(&) by weight	
Adflex [®] Q 200 F	79.4
Sunpar [®] 2280	20
Irganox [®] PS 802	0.4
Irganox [®] 1010	0.2

The process used for manufacturing the cable was the following.

E1030

31

The Adflex[®] Q 200 F was fed directly into the extruder hopper. An extruder having a diameter of 80 mm and an L/D ratio of 25 was used. Subsequently, an attempt was made to inject the Sunpar[®] 2280 previously mixed with the antioxidants into the extruder. The injection was impossible to be carried out since the dielectric liquid exit the extruder die. Consequently, the production of a finished cable was impossible to be carried out.

10

15

20

25

30

35

C1030

32

CLAIMS

1. Cable comprising at least one electrical conductor and at least one extruded covering layer based on a thermoplastic polymer material in admixture with a dielectric liquid, wherein:
- 5 - said thermoplastic polymer material is selected from:
- (a) at least one propylene homopolymer or at least one copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 130°C and a melting enthalpy of from 20 J/g to 100 J/g;
- 10 (b) a mechanical mixture comprising at least one propylene homopolymer or copolymer (a) and (c) at least one elastomeric copolymer of ethylene with at least one aliphatic α -olefin, and optionally a polyene;
- 15 - the concentration by weight of said dielectric liquid in said thermoplastic polymer material is less or equal to the saturation concentration of said dielectric liquid in said thermoplastic polymer material;
- 20 - said dielectric liquid has the following characteristics:
- 25 - an amount of polar compounds lower than or equal to 2.5% by weight with respect to the total weight of the dielectric liquid;
- 30 - a melting point or a pour point lower than 80°C;
- 35

C1030

33

- a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms lower than 0.6, when the dielectric liquid is aromatic.

- 5 2. Cable according to claim 1, wherein the propylene homopolymer or copolymer (a) has a melting point of from 140°C to 170°C.
- 10 3. Cable according to claim 1 or 2, wherein the propylene homopolymer or copolymer (a) has a melting enthalpy of from 30 J/g to 85 J/g.
- 15 4. Cable according to any one of the preceding claims, wherein the propylene homopolymer or copolymer (a) has a flexural modulus, measured according to ASTM standard D790, at room temperature, of from 30 MPa to 1400 MPa.
- 20 5. Cable according to claim 4, wherein the propylene homopolymer or copolymer (a) has a flexural modulus, measured according to ASTM standard D790, at room temperature, of from 60 MPa to 1000 MPa.
- 25 6. Cable according to any one of the preceding claims, wherein the propylene homopolymer or copolymer (a) has a melt flow index (MFI), measured at 230°C with a load of 21.6 N according to ASTM standard D1238/L, of from 0.05 dg/min to 10.0 dg/min.
- 30 7. Cable according to claim 6, wherein the propylene homopolymer or copolymer (a) has a melt flow index (MFI), measured at 230°C with a load of 21.6 N according to ASTM standard D1238/L, of from 0.4 dg/min to 5.0 dg/min.
- 35 8. Cable according to any one of the preceding claims, wherein in the propylene copolymer (a) the olefin comonomer is present in a quantity of less than or equal to 15 mol%.
9. Cable according to claim 8, wherein in the

propylene copolymer (a) the olefin comonomer is present in a quantity of less than or equal to 10 mol%,

- 5 10. Cable according to any one of the preceding claims, wherein in the propylene copolymer (a) the olefin comonomer is ethylene or an α -olefin of formula $\text{CH}_2=\text{CH}-\text{R}$, where R is a linear or branched C_2-C_{10} alkyl.
- 10 11. Cable according to claim 10, wherein the α -olefin is selected from: 1-butene, 1-pentene, 4-methyl-1-pentene, 1-hexene, 1-octene, 1-decene, 1-dodecene, or mixtures thereof.
- 15 12. Cable according to any one of the preceding claims, wherein the propylene homopolymer or copolymer (a) is selected from:
- 20 (a₁) a propylene homopolymer or a copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, having a flexural modulus generally of from 30 MPa to 900 MPa;
- 25 (a₂) a heterophase copolymer comprising a thermoplastic phase based on propylene and an elastomeric phase based on ethylene copolymerized with an α -olefin.
- 30 13. Cable according to claim 12, wherein the propylene homopolymer or copolymer of class (a₁) has a melting point of from 140°C to 170°C.
14. Cable according to claim 12 or 13, wherein the propylene homopolymer or copolymer of class (a₁) has a melting enthalpy of from 30 J/g to 80 J/g.
- 35 15. Cable according to any one of claims 12 to 14, wherein the propylene homopolymer or copolymer of class (a₁) has a fraction soluble in boiling diethyl ether in an amount of less than or equal to 12 wt%, having a melting enthalpy of less

PC1030

35

than or equal to 4 J/g.

16. Cable according to any one of claims 12 to 15,
wherein the propylene homopolymer or copolymer
of class (a₁) has a fraction soluble in boiling
n-heptane in an amount of from 15 wt% to 60 wt%,
having a melting enthalpy of from 10 J/g to 40
J/g.
17. Cable according to any one of claims 12 to 16,
wherein the propylene homopolymer or copolymer
of class (a₁) has a fraction insoluble in boiling
n-heptane in an amount of from 40 wt% to 85 wt%,
having a melting enthalpy of greater than or
equal to 45 J/g.
18. Cable according to claim 12, wherein the α-
olefin included in the elastomeric phase of a
heterophase copolymer of class (a₂) is propylene.
19. Cable according to claim 12, wherein the
heterophase copolymer of class (a₂) is a
heterophase copolymer in which the elastomeric
phase consists of an elastomeric copolymer of
ethylene and propylene comprising from 15 wt% to
50 wt% of ethylene and from 50 wt% to 85 wt% of
propylene with respect to the weight of the
elastomeric phase.
20. Cable according to any one of the preceding
claims, wherein the elastomeric copolymer of
ethylene (c) has a melting enthalpy of less than
30 J/g.
21. Cable according to any one of the preceding
claims, wherein the quantity of the elastomeric
copolymer (c) is less than 70% with respect to
the total weight of the thermoplastic base
material.
22. Cable according to any one of the preceding
claims, wherein in the elastomeric copolymer of
ethylene (c), the term "aliphatic α-olefin" is

21030

36

an olefin of formula $\text{CH}_2=\text{CH-R}$, in which R represents a linear or branched alkyl group containing from 1 to 12 carbon atoms.

23. Cable according to claim 22, wherein the
5 aliphatic α -olefin is selected from propylene, 1-butene, isobutylene, 1-pentene, 4-methyl-1-pentene, 1-hexene, 1-octene, 1-dodecene, or mixtures thereof.
24. Cable according to claim 23, wherein the
10 aliphatic α -olefin is selected from propylene, 1-hexene, 1-octene.
25. Cable according to any one of the preceding claims, wherein in the elastomeric copolymer of ethylene (c), the term "polyene" is a conjugated
15 or non-conjugated diene, triene or tetraene.
26. Cable according to claim 25, wherein the polyene is a diene.
27. Cable according to any one of the preceding claims, wherein the elastomeric copolymer of
20 ethylene (c) is selected from:
(c₁) copolymers having the following monomer composition: 35 mol%-90 mol% of ethylene; 10 mol%-65 mol% of an aliphatic α -olefin; 0 mol%-10 mol% of a polyene.
- 25 (c₂) copolymers having the following monomer composition: 75 mol%-97 mol% of ethylene; 3 mol%-25 mol% of an aliphatic α -olefin; 0 mol%-5 mol% of a polyene.
28. Cable according to any one of the preceding
30 claims, wherein the dielectric liquid has an amount of polar compounds of between 0.1 and 2.3.
29. Cable according to any one of the preceding
35 claims, wherein the dielectric liquid has a melting point or a pour point of between -130°C and +80°C.

1030

37

30. Cable according to any one of the preceding claims, wherein the dielectric liquid has a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms of between 0.01 and 0.4.
31. Cable according to any one of the preceding claims, wherein the dielectric liquid has a dielectric constant, at 25°C, of less than or equal to 3.5 (measured in accordance with IEC 247).
32. Cable according to any one of the preceding claims, wherein the dielectric liquid has a viscosity, at 40°C, of between 10 cSt and 800 cSt (measured according to ASTM standard D445).
33. Cable according to claim 32, wherein the dielectric liquid has a viscosity, at 40°C, of between 20 cSt and 500 cSt (measured according to ASTM standard D445).
34. Cable according to any one of the preceding claims, wherein the dielectric liquid is selected from: mineral oils such as naphthenic oils, aromatic oils, paraffinic oils, polyaromatic oils, said mineral oils optionally containing at least one heteroatom selected from oxygen, nitrogen or sulphur; liquid paraffins; vegetable oils such as soybean oil, linseed oil, castor oil; oligomeric aromatic polyolefins; paraffinic waxes such as polyethylene waxes, polypropylene waxes; synthetic oils such as silicone oils, alkyl benzenes such as dodecylbenzene, di(octylbenzyl)toluene, aliphatic esters such as tetraesters of pentaerythritol, esters of sebacic acid, phthalic acid esters, olefin oligomers such as optionally hydrogenated polybutenes or polyisobutenes; or mixtures thereof.

C1030

38

35. Cable according to claim 34, wherein the dielectric liquid is selected from paraffinic oils, naphthenic oils.
- 5 36. Cable according to any one of the preceding claims, wherein the weight ratio of dielectric liquid to thermoplastic polymer material is between 1:99 and 25:75.
- 10 37. Cable according to claim 36, wherein the weight ratio of dielectric liquid to thermoplastic polymer material is between 2:98 and 20:80.
38. Cable according to claim 37, wherein the weight ratio of dielectric liquid to thermoplastic polymer material is between 3:97 and 10:90.
- 15 39. Cable according to any one of the preceding claims, wherein the thermoplastic polymer material is selected from propylene homopolymers or copolymers comprising at least 40 wt% of amorphous phase, with respect to the total polymer weight.
- 20 40. Cable according to any one of the preceding claims, wherein the extruded covering layer based on said thermoplastic polymer material in admixture with said dielectric liquid is an electrically insulating layer.
- 25 41. Cable according to any one of the preceding claims, wherein the extruded covering layer based on said thermoplastic polymer material in admixture with said dielectric liquid is a semiconductive layer.
- 30 42. Polymer composition comprising a thermoplastic polymer material in admixture with a dielectric liquid, wherein:
- said thermoplastic polymer material is selected from:

35 (a) at least one propylene homopolymer or at least one copolymer of propylene with at

C1030

39

least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 130°C and a melting enthalpy of from 20 J/g to 100 J/g;

(b) a mechanical mixture comprising at least one propylene homopolymer or copolymer (a) and (c) at least one elastomeric copolymer of ethylene with at least one aliphatic α -olefin, and optionally a polyene;

- the concentration by weight of said dielectric liquid in said thermoplastic polymer material is less or equal to the saturation concentration of said dielectric liquid in said thermoplastic polymer material;

- said dielectric liquid has the following characteristics:

- an amount of polar compounds lower than or equal to 2.5% by weight with respect to the total weight of the dielectric liquid;

- a melting point or a pour point lower than 80°C;

- a ratio of number of aromatic carbon atoms with respect to the total number of carbon atoms lower than 0.6, when the dielectric liquid is aromatic.

43. Polymer composition according to claim 42, wherein the thermoplastic polymer material is defined according to any one of claims 2 to 27.

44. Polymer composition according to claim 42 or 43, wherein the dielectric liquid is defined according to any one of claims 28 to 38.

1030

40

45. Use of the polymer composition defined according to any one of claim 42 to 44, as the polymer base material for preparing a cable covering layer with electrical insulation properties.

5 45. Use of the polymer composition defined according to any one of claim 42 to 44, as the polymer base material for preparing a cable covering layer with semiconductive properties.

10

15

20

25

30

35

ABSTRACT

Cable comprising at least one electrical conductor and at least one extruded covering layer based on a thermoplastic polymer material in admixture with a dielectric liquid, wherein:

- said thermoplastic polymer material is selected from:

(a) at least one propylene homopolymer or at least one copolymer of propylene with at least one olefin comonomer selected from ethylene and an α -olefin other than propylene, said homopolymer or copolymer having a melting point greater than or equal to 130°C and a melting enthalpy of from 20 J/g to 100 J/g;

(b) a mechanical mixture comprising at least one propylene homopolymer or copolymer (a) and (c) at least one elastomeric copolymer of ethylene with at least one aliphatic α -olefin, and optionally a polyene;

- the concentration by weight of said dielectric liquid in said thermoplastic polymer material is less or equal to the saturation concentration of said dielectric liquid in said thermoplastic polymer material;

- said dielectric liquid has the following characteristics:

- an amount of polar compounds lower than or equal to 2.5% by weight with respect to the total weight of the dielectric liquid;

- a melting point or a pour point lower than 80°C;

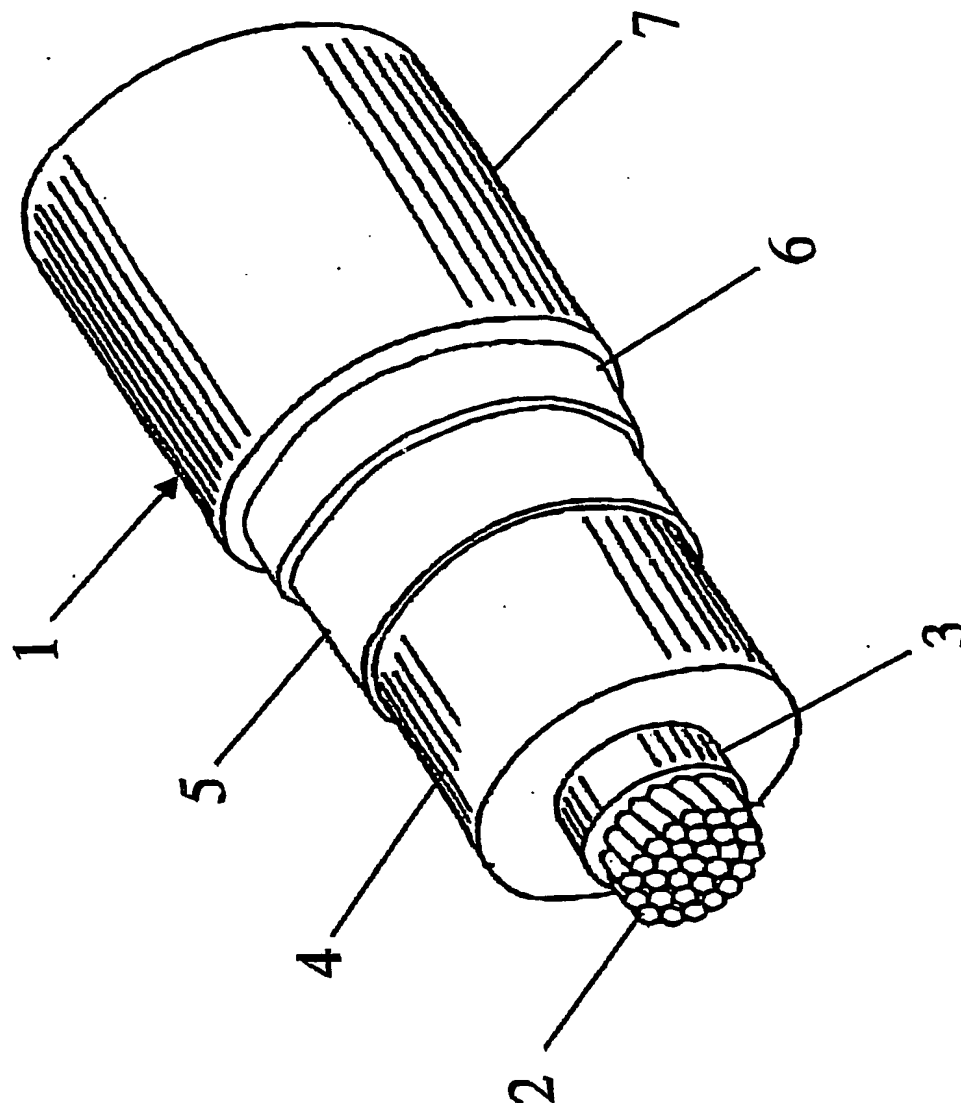
- a ratio of number of aromatic carbon atoms with respect to the total number of carbon

atoms lower than 0.6, when the dielectric liquid is aromatic.

5 The cable of the invention possesses good mechanical and electrical properties, including high dielectric strength, in particular enabling it to be used at high operating temperature.

Fig. 1.

Fig. 1



**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

☐ BLACK BORDERS

☒ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES

☐ FADED TEXT OR DRAWING

☐ BLURRED OR ILLEGIBLE TEXT OR DRAWING

☐ SKEWED/SLANTED IMAGES

☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS

☐ GRAY SCALE DOCUMENTS

☐ LINES OR MARKS ON ORIGINAL DOCUMENT

☒ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY

☐ OTHER: _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.